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PEF A

BULLVEL

MEMORANDUM

February 9, 1999

SUBJECT: Atrazine Qualitative Risk Assessment Based On Female

Sprague-Dawley Rat Dietary Study

P.C. Code 080803

TO:

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Background

A carcinogenicity toxicity study in female Sprague-Dawley rats was conducted by Covance Laboratories, Vienna, Virginia, for Novartis Crop Protection, Greensboro, North Carolina, and dated April 15, 1998 (Lab Report No. 2386-108; MRID No. 445447-01).

The study design allocated groups of 160 female rats to dose levels of 0, 25, 50, 70, or 400 ppm of Atrazine for 105 weeks. Eighty animals per dose group were ovariectomized while the others were left intact. Only the intact animals have been included in this analysis, as there were no compound-related tumors observed in the ovariectomized animals.

<u>Survival Analyses</u>

The statistical evaluation of mortality indicated a significant <u>increasing</u> trend with increasing doses of Atrazine in female rats. See Table 1 for mortality test results.

The statistical evaluation of mortality was based upon the Thomas, Breslow and Gart computer program.

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Tumor Analyses

Female rats had significant increasing trends, and significant differences in the pair-wise comparisons of the 400 ppm dose group with the controls, for mammary gland carcinomas and fibroadenomas and/or adenomas and/or carcinomas combined, all at p < 0.01. There were significant differences in the pair-wise comparisons of the 25 and 70 ppm dose groups with the controls for mammary gland fibroadenomas and fibroadenomas and/or adenomas and/or carcinomas combined, all at p < 0.05. There were significant differences in the pair-wise comparison of the 50 ppm dose group with the controls for mammary gland fibroadenomas and fibroadenomas and/or adenomas and/or carcinomas combined, both at p < 0.01. There was also a significant difference in the pair-wise comparison of the 400 ppm dose group with the controls for mammary gland fibroadenomas at p < 0.05.

The statistical analyses of the female rats were based upon Peto's Prevalence Test since there was a statistically significant positive trend for mortality with increasing doses of Atrazine in female rats. See Table 2 for tumor analysis results.

Table 1. Atrazine Sprague-Dawley <u>Female</u> Rat Study
Intact Animals ONLY

Mortality Rates and Cox or Generalized K/W Test Results

			<u>Weeks</u>			
Dose (ppm)	1-26	27-52	53 ⁱ	53-78	79-106 ^f	Total
0	1/80	1/79	20/78	9/58	23/49	34/60 (57)*
25	0/80	1/80	21/79	14/58	25/44	40/5 ₉ 9 (68)
50	1/80	2/79	21/77	12/56	27/44	42/59 (71)
70	0/80	2/80	20/78	8/58	31/50	41/60 (68)
400	2/80	1/78	22/77	16/55	26/39	45/58 (78)*

'Number of animals that died during interval/Number of animals alive at the beginning of the interval.

() Percent.

Note: Time intervals were selected for display purposes only.

Significance of trend denoted at <u>control</u>.

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fFinal sacrifice at week 105.

ⁱInterim sacrifice at week 53.

Significance of pair-wise comparison with control denoted at dose level.

If * , then p < 0.05. If ** , then p < 0.01.

Table 2. Atrazine Sprague-Dawley <u>Female</u> Rat Study
Intact Animals ONLY

Mammary Gland Tumor Rates and Peto's Prevalence Test Results (p values)

		Dose (ppm)						
	0	25	50	70	400			
Fibro- adenomas (%)	16/78 (21)	25ª/79 (32)	34/77 (44)	29/78 (37)	25/77 (32)			
p =	0.233	0.030*	0.000**	0.014*	0.014*			
Adenomas (%)	0/28	0/24 (0)	1 ^b /20 (5)	0/21 (0)	0/15			
Carcinomas (%)	12/80 (15)	18/80 (22)	20/79 (25)	14/80 (18)	27°/80 (34)			
p =	0.002**	0.112	0.067	0.395	0.007**			
Combined ^d (%)	24/80 (30)	34/80 (42)	45/79 (57)	38/80 (48)	43/80 (54)			
p =	0.004**	0.025*	0.000**	0.015*	0.000**			

*Number of tumor bearing animals/Number of animals examined, excluding those that died or were sacrificed before observation of the first tumor.

 $^{
m d}$ The number of animals with multiple tumors are 4, 9, 10, 5 and 9 for the control, 25, 50, 70 and 400 ppm dose groups, respectively.

Note: Significance of trend denoted at control.

Significance of pair-wise comparison with control denoted at dose level.

^{*}First fibroadenoma observed at week 53, dose 25 ppm.

bFirst adenoma observed at week 103, dose 50 ppm.

[°]First carcinoma observed at week 19, dose 400 ppm.

If *, then p < 0.05. If **, then p < 0.01.

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References

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